WHAT IS CLAIMED IS:

1. A protein having a $\beta 1,3$ -galactosyltransferase activity derived from a microorganism having an activity of transferring galactose to N-acetylglucosamine with $\beta 1,3$ -linkage.

- 2. The protein according to claim 1, wherein the microorganism belongs to the genus Streptococcus.
- 3. The protein according to claim 2, wherein the microorganism is Streptococcus agalactiae.

A protein comprising the amino acid sequence represented by SEQ ID NO:1.

5. A protein comprising an amino acid sequence in which at most 20 amino acids are deleted, replaced, inserted or added in the amino acid sequence represented by SEQ ID NO: , said protein having a β 1,3-galactosyltrans erase activity.

DNA encoding the protein of any one of claims 1 to 5.

A DNA comprising the nucleotide sequence represented by SEQ ID NO:2.

8. A DNA which hybridizes with a DNA comprising the complementary sequence to the nucleotide sequence represented by SEQ ID NO:2 under stringent conditions, and encodes a protein having a $\beta1,3$ -galactosyltransferase activity.

9. A recombinant DNA comprising the DNA of any one of claims 6 to 8 and a vector.

- 10. A transformant obtained by introducing the recombinant DNA of claim 9 into a host cell.
- 11. The transformant according to claim 10, wherein the host cell is a microorganism.
- 12. The transformant according to claim 11, wherein the microgranism belongs to the genus Escherichia.
- 13. The transformant according to claim 12, wherein the microorganism belonging to the genus Escherichia is Escherichia coli.

 β 14. A method for producing a protein having a β 1,3-galactosyltransferase activity, comprising:

culturing the transformant of any one of claims 10 to 13 in a medium to produce and accumulate a protein having a $\beta 1,3$ -galactosyltransferase activity in the culture, and

recovering the protein from the culture.

15. A method for producing a galactose-containing carbohydrate, comprising:

selecting, as an enzyme source, a culture of the transformant of any one of claims 10 to 13 or a treated product of the culture,

allowing the enzyme source, uridine-5'diphosphogalactose and an acceptor carbohydrate to be
present in an aqueous medium to produce and accumulate the
galactose-containing carbohydrate in the aqueous medium,
and

recovering the galactose-containing carbohydrate from the aqueous medium.

16. The method according to claim 15, wherein the treated product of the culture is selected from the group consisting of a concentrated product of the culture, a dried product of the culture, cells obtained by centrifuging the culture, a dried product of the cells, a

freeze-dried product of the cells, a surfactant-treated product of the cells, an ultrasonic-treated product of the cells, a mechanically disrupted product of the cells, a solvent-treated product of the cells, an enzyme-treated product of the cells, an immobilized product of the cells and an enzyme preparation obtained by extracting from the cells.

- 17. The method according to claim 15, wherein the acceptor carbohydrate is a carbohydrate having N-acetylglucosamine at its non-reducing terminal.
- 18. The method according to claim 15, wherein the acceptor carbohydrate is selected from the group consisting of N-acetylglucosamine and lacto-N-triose II.
- 19. The method according to claim 15, wherein the galactose-containing carbohydrate is selected from the group consisting of lacto-N-biose and lacto-N-tetraose.

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